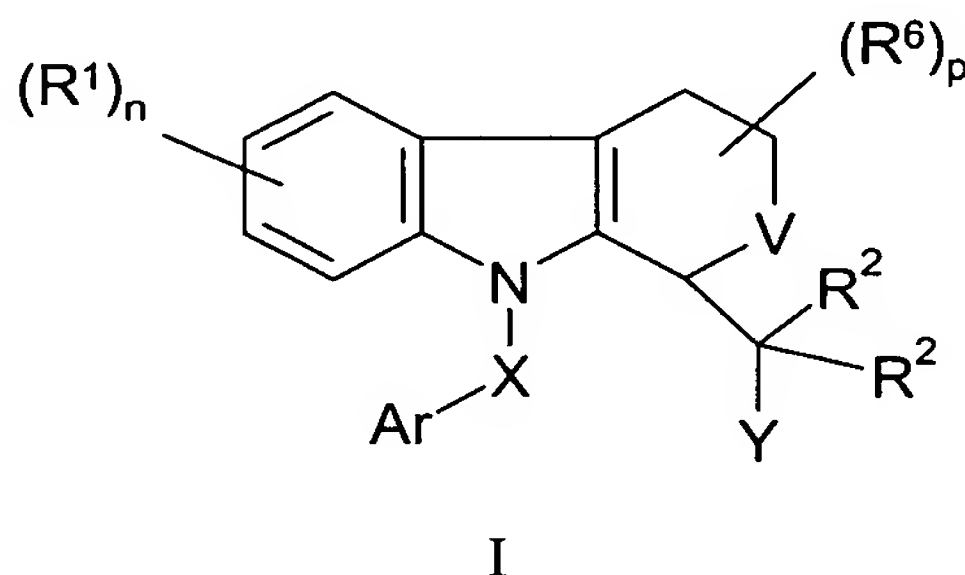


Amendments to the Claims:

The listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims

Claim 1 (Currently amended) ~~The use, for the manufacture of a medicament for treatment or prevention of~~ A method of treating or preventing a disease associated with the deposition of β -amyloid in the brain, comprising administering to a patient in need thereof a therapeutically effective amount of a compound of formula I:



wherein V represents a bond, CH₂ or CH₂CH₂;

X represents SO₂ or CHR³ where R³ is H or a hydrocarbon group containing up to 10 carbon atoms which is optionally substituted with halogen, CF₃, C₁₋₄alkoxy or C₁₋₄alkylthio;

Y represents CO₂H or tetrazole;

Ar represents phenyl which optionally bears up to 3 substituents independently selected from hydrocarbon groups of up to 6 carbon atoms and (CH₂)_m-Z where m is 0, 1 or 2 and Z represents halogen, N₃, CN, CF₃, OCF₃, OR⁴, S(O)_tR⁴ where t is 0, 1 or 2, CO₂R⁴, tetrazole, N(R⁴)₂, NHCOR⁵, NHCON(R⁴)₂, CON(R⁴)₂, SO₂N(R⁴)₂, NHSO₂R⁵, COR⁵, or OCOR⁵;

n is 0, 1, 2 or 3;

each R¹ is independently selected from nonaromatic hydrocarbon groups of up to 6 carbon atoms and (CH₂)_q-W where q is 0, 1 or 2 and W represents halogen, CN, CF₃, OR⁴, N(R⁴)₂, S(O)_tR⁴ where t is 0, 1 or 2, CO₂R⁴, tetrazole, CON(R⁴)₂, SO₂N(R⁴)₂, COR⁵, OCOR⁵ or phenyl or heteroaryl either of which optionally bears up to 3 substituents selected from halogen, CF₃, OCF₃, CN, OH, C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkylthio or C₁₋₄alkoxycarbonyl;

each R² is independently H or C₁₋₄alkyl; or one R² group together with an R⁶ group attached at the same ring position as the -C(R²)₂-Y moiety completes a spiro-linked hydrocarbon ring of 3-6 members;

R⁴ represents H or a hydrocarbon group of up to 7 carbon atoms, optionally substituted with halogen, CN, CF₃, OH, C₁₋₄alkoxy or C₁₋₄alkoxycarbonyl; or two R⁴ groups attached to the same nitrogen atom may complete a 5- or 6-membered heterocyclic ring;

R^5 represents R^4 that is other than H;

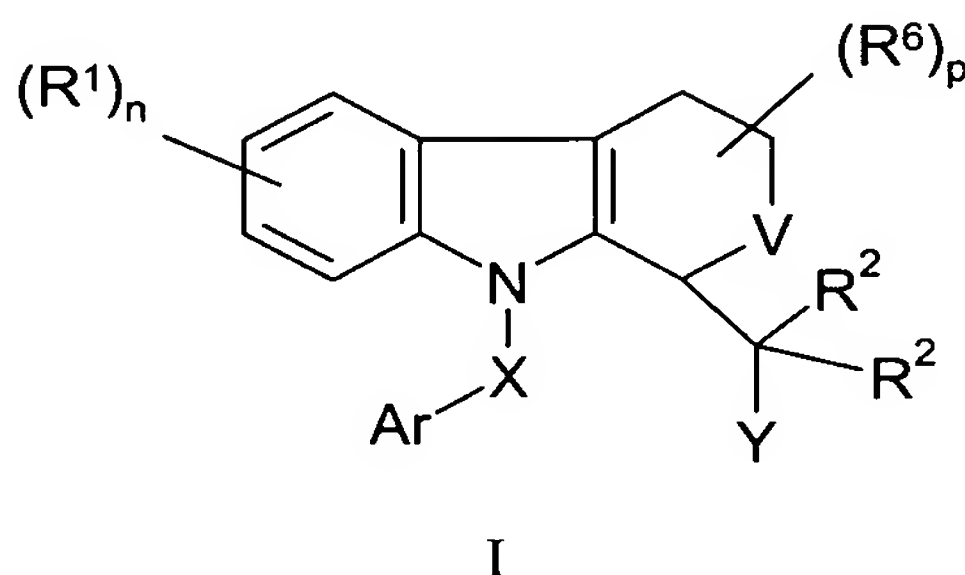
p is 0, 1 or 2; and

R^6 represents C_{1-6} alkyl, C_{2-6} alkenyl or phenyl, benzyl or heteroaryl, said phenyl, benzyl or heteroaryl optionally bearing up to 3 substituents selected from halogen, CN, CF_3 , OCF_3 , OR^4 , CO_2R^4 , COR^5 , $OCOR^5$ and C_{1-4} alkyl; or an R^6 group together with an R^2 group may complete a spiro-linked hydrocarbon ring as defined previously; or a pharmaceutically acceptable salt thereof.

Claim 2 (Cancelled)

Claim 3 (Currently amended) ~~Use according to~~ The method of claim 1 wherein said disease is Alzheimer's disease, cerebral amyloid angiopathy, multi-infarct dementia, dementia pugilistica or Down syndrome.

Claim 4 (Currently amended) A compound according to formula I ~~as defined in claim 1~~ wherein



wherein V represents a bond, CH_2 or CH_2CH_2 ;

X represents SO_2 or CHR^3 where R^3 is H or a hydrocarbon group containing up to 10 carbon atoms which is optionally substituted with halogen, CF_3 , C_{1-4} alkoxy or C_{1-4} alkylthio;

Y represents CO_2H or tetrazole;

Ar represents phenyl which optionally bears up to 3 substituents independently selected from hydrocarbon groups of up to 6 carbon atoms and $(CH_2)_m-Z$ where m is 0, 1 or 2 and Z represents halogen, N_3 , CN, CF_3 , OCF_3 , OR^4 , $S(O)_tR^4$ where t is 0, 1 or 2, CO_2R^4 , tetrazole, $N(R^4)_2$, $NHCOR^5$, $NHCON(R^4)_2$, $CON(R^4)_2$, $SO_2N(R^4)_2$, $NHSO_2R^5$, COR^5 , or $OCOR^5$;

n is 0, 1, 2 or 3;

each R^1 is independently selected from nonaromatic hydrocarbon groups of up to 6 carbon atoms and $(CH_2)_q-W$ where q is 0, 1 or 2 and W represents halogen, CN, CF_3 , OR^4 , $N(R^4)_2$, $S(O)_tR^4$ where t is 0, 1 or 2, CO_2R^4 , tetrazole, $CON(R^4)_2$, $SO_2N(R^4)_2$, COR^5 , $OCOR^5$ or phenyl or

heteroaryl either of which optionally bears up to 3 substituents selected from halogen, CF₃, OCF₃, CN, OH, C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkylthio or C₁₋₄alkoxycarbonyl;

each R² is independently H or C₁₋₄alkyl; or one R² group together with an R⁶ group attached at the same ring position as the -C(R²)₂-Y moiety completes a spiro-linked hydrocarbon ring of 3-6 members;

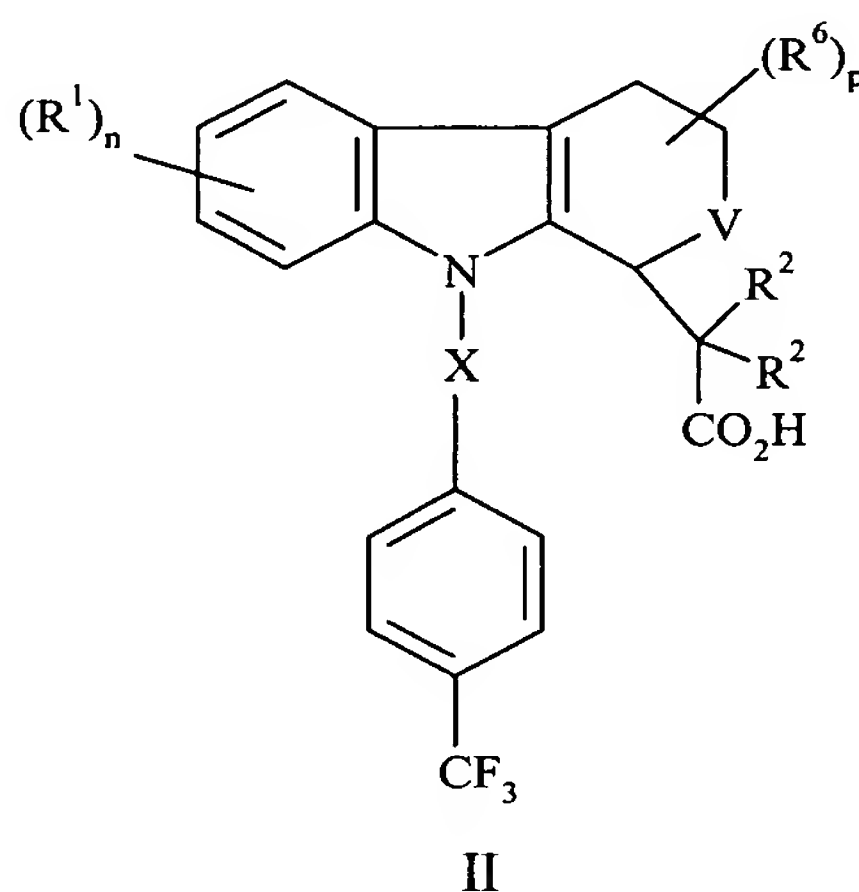
R⁴ represents H or a hydrocarbon group of up to 7 carbon atoms, optionally substituted with halogen, CN, CF₃, OH, C₁₋₄alkoxy or C₁₋₄alkoxycarbonyl; or two R⁴ groups attached to the same nitrogen atom may complete a 5- or 6-membered heterocyclic ring;

R⁵ represents R⁴ that is other than H;

p is 1 or 2;

and at least one R⁶ represents C₂₋₆ alkenyl or optionally-substituted phenyl, heteroaryl or benzyl; or a pharmaceutically acceptable salt thereof.

Claim 5 (Currently amended) A compound according to formula II:

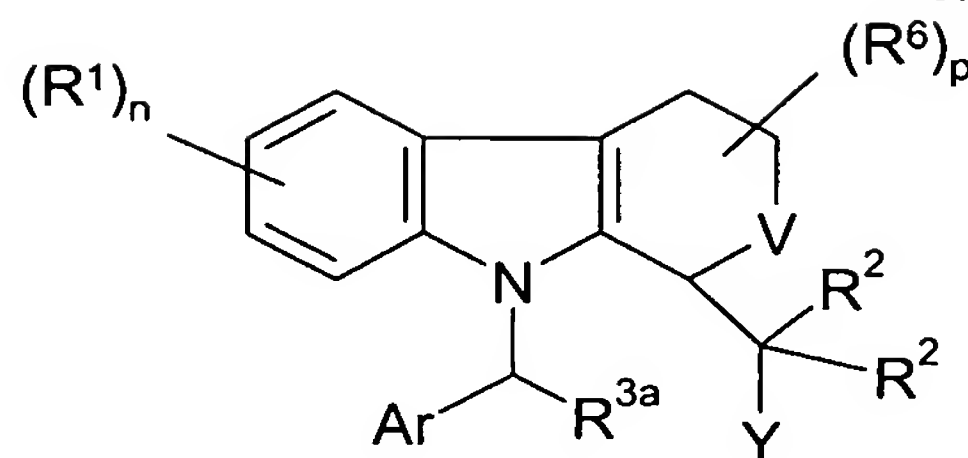


or a pharmaceutically acceptable salt thereof, where V, X, n, p, R¹, R² and R⁶ are as defined in claim 4;

with the proviso that if V is CH₂, X is CH₂, p is zero and each R² is H, then (R¹)_n does not represent 6,8-difluoro.

Claim 6 (Currently amended) A compound according to claim 4 ~~or claim 5~~ wherein X is CHR³.

Claim 7 (Currently amended) A compound according to formula III:



III

or a pharmaceutically acceptable salt thereof, wherein R^{3a} represents a hydrocarbon group containing from 2 to 10 carbon atoms which is optionally substituted with halogen, CF₃, C₁₋₄alkoxy or C₁₋₄alkylthio; and the remaining variables are as defined in claim 4, with the proviso that R¹ does not represent SOR⁴ or SO₂R⁴.

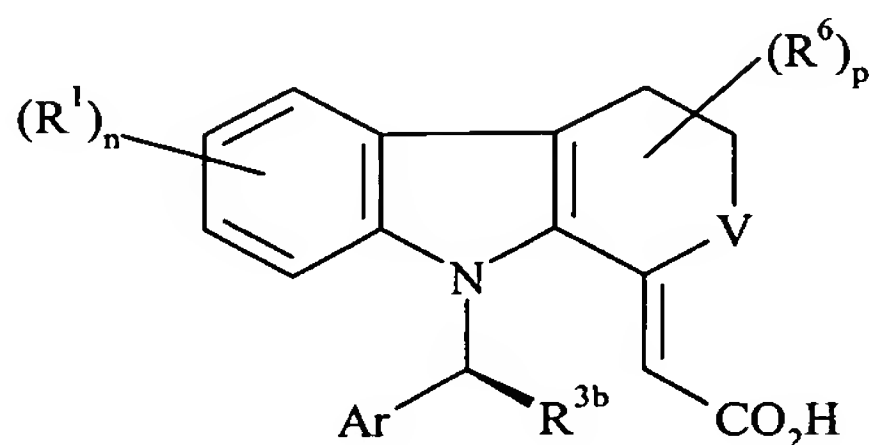
Claim 8 (Original) A compound according to claim 7 wherein Y represents CO₂H, Ar represents 4-trifluoromethylphenyl, and both R² groups represent H.

Claim 9 (Currently amended) A compound according to ~~any of claims 4-8~~ claim 4 wherein n is 1 or 2 and each R¹ is independently selected from methyl, ethyl, isopropyl, n-butyl, t-butyl, cyclopropyl, Br, Cl, F, CN, CF₃, OCH₃, OCF₃, SCH₃, morpholin-1-yl, 4-fluorophenyl, 3,4-dichlorophenyl, 3-methylthiophenyl, 2,5-dimethylphenyl and 3-trifluoromethoxyphenyl.

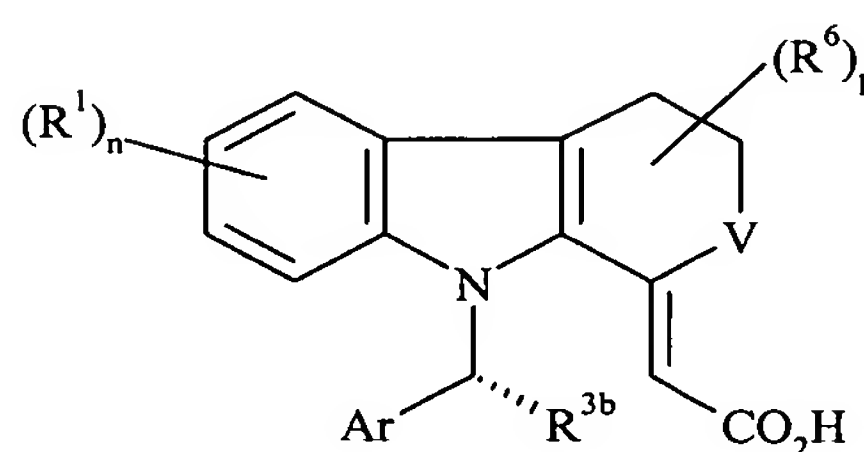
Claim 10 (Cancelled)

Claim 11 (Currently amended) A pharmaceutical composition comprising a compound according to ~~any of claims 4-9~~ claim 4 and a pharmaceutically acceptable carrier.

Claim 12 (Original) A process for preparing a compound of formula III as defined in claim 7 comprising the step of hydrogenating a compound of formula (11a) or (11b) over a chiral Ru(BINAP)Cl₂ catalyst:



(11a)



(11b)

wherein BINAP is bis(diphenylphosphino)-1,1'-binaphthyl and R^{3b} is R^3 that is other than H.

Claim 13 (Original) The process of claim 12 wherein the compound of formula (11a) or (11b) is obtained by reaction of a compound of formula (5a) or (5b) with a compound of formula (10):

